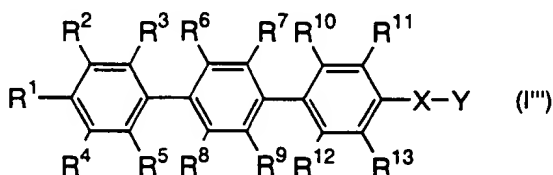


Claim 14 (Amended)

Line 1, after "the formula (I)" insert --according to claim
13--

Line 2, delete "according to claim 13"

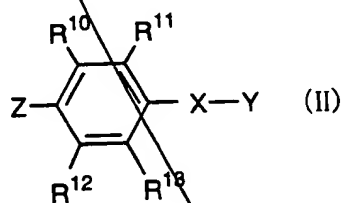
Claim 15 (Amended) A process for producing a compound of the
formula (I'''):



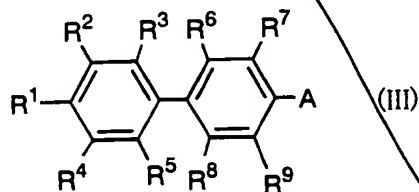
wherein R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸, R⁹, R¹⁰, R¹¹, R¹² and R¹³ are each independently hydrogen, hydroxy, halogen, carboxy, optionally substituted lower alkyl, optionally substituted lower alkoxy, optionally substituted lower alkenyl, optionally substituted lower alkenyloxy, optionally substituted lower alkylthio, optionally substituted lower alkoxycarbonyl, optionally substituted acyloxy, optionally substituted lower alkylsulfonyl, optionally substituted lower alkylsulfonyloxy, optionally substituted lower alkylsulfinyl, nitro, cyano, formyl, optionally substituted amino, optionally substituted carbamoyl, optionally substituted sulfamoyl or optionally substituted heterocyclyl,

~~X is -O-, -CH₂-, NR¹⁴- wherein R¹⁴ is hydrogen, optionally substituted lower alkyl, optionally substituted lower alkenyl or acetyl, or -S(O)_p- wherein p is an integer of 0 to 2, Y is optionally substituted lower alkyl, optionally substituted lower alkenyl, optionally substituted lower alkynyl, optionally substituted acyl, optionally substituted cycloalkyl, optionally substituted cycloalkenyl, optionally substituted aryl or optionally substituted heterocyclyl, and Y may optionally be substituted lower alkoxy when X is -CH₂- and may optionally be substituted lower alkoxycarbonyl, optionally substituted lower alkylsulfonyl or optionally substituted arylsulfonyl when X is -O- or -NR¹⁴, R¹ and R⁴, R¹ and R², R² and R³, R⁴ and R⁵, R⁶ and R⁷, R⁸ and R⁹, R¹⁰ and R¹¹, R¹² and R¹³, R¹¹ and -X-Y, or R¹³ and -X-Y taken together may form a 5- or 6-membered ring which may contain one or more of O, S or NR¹⁵ wherein R¹⁵ is hydrogen, optionally substituted lower alkyl, optionally substituted lower alkenyl, optionally substituted arylsulfonyl, and which may optionally be substituted, excluding a compound wherein one or more of R⁶, R⁷, R⁸ and R⁹ are halogen and the others are hydrogen, compounds wherein all of R⁶, R⁷, R⁸ and R⁹ are halogen and compounds wherein all of R²-R¹³ are hydrogen, halogen or cyano, provided that R¹ is not hydrogen, fluorine, optionally substituted lower alkyl or optionally substituted lower alkoxy, all of R², R³, R⁴, R⁵ and R¹²~~

are hydrogen or R^{13} is not hydrogen or halogen when R^6 , R^7 , R^8 and R^9 are all simultaneously hydrogen, and further provided that R^1 is not methyl or acetyloxy, R^{13} is not hydrogen, optionally substituted lower alkoxy, carbonyl or optionally substituted carbamoyl or $-X-Y$ is not methoxy when at least one of R^6 , R^7 , R^8 and R^9 is a substituent other than hydrogen, pharmaceutically acceptable salt or hydrate thereof, which comprises reacting a compound of the formula (II):



with a compound of the formula (III):



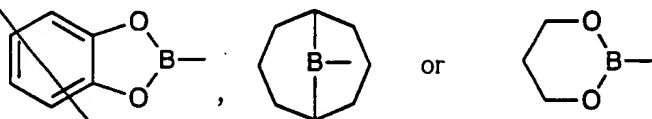
wherein, in the formulas (II) and (III), [R^1-R^{13} , X and Y are the same as defined in claim 7,] R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 , R^9 , R^{10} , R^{11} , R^{12} and R^{13} are each independently hydrogen, hydroxy, halogen, carboxy, optionally substituted lower alkyl, optionally substituted lower alkoxy, optionally substituted lower alkenyl, optionally substituted lower alkenyloxy, optionally substituted

lower alkylthio, optionally substituted lower alkoxy
carbonyl, optionally substituted acyloxy, optionally substituted lower
alkylsulfonyl, optionally substituted lower alkylsulfonyloxy,
optionally substituted lower alkylsulfinyl, nitro, cyano, formyl,
optionally substituted amino, optionally substituted carbamoyl,
optionally substituted sulfamoyl or optionally substituted
heterocyclyl,

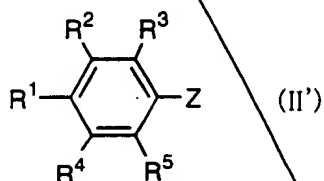
X is -O-, -CH₂-, NR¹⁴- wherein R¹⁴ is hydrogen, optionally
substituted lower alkyl, optionally substituted lower alkenyl or
acetyl, or -S(O)_p- wherein p is an integer of 0 to 2, Y is
optionally substituted lower alkyl, optionally substituted lower
alkenyl, optionally substituted lower alkynyl, optionally
substituted acyl, optionally substituted cycloalkyl, optionally
substituted cycloalkenyl, optionally substituted aryl or
optionally substituted heterocyclyl, and Y may optionally be
substituted lower alkoxy when X is -CH₂- and may optionally be
substituted lower alkoxy, optionally substituted lower
alkylsulfonyl or optionally substituted arylsulfonyl when X is -O-
or -NR¹⁴, R¹ and R⁴, R¹ and R², R² and R³, R⁴ and R⁵, R⁶ and R⁷, R⁸
and R⁹, R¹⁰ and R¹¹, R¹² and R¹³, R¹¹ and -X-Y, or R¹³ and -X-Y taken
together may form a 5- or 6-membered ring which may contain one or
more of O, S or NR¹⁵ wherein R¹⁵ is hydrogen, optionally
substituted lower alkyl, optionally substituted lower alkenyl,

optionally substituted arylsulfonyl, and which may optionally be substituted,

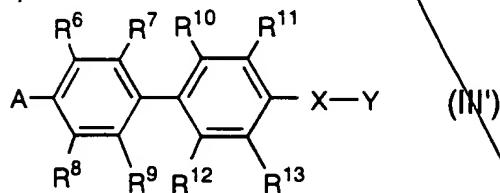
either of A and Z is dihydroxyborane, di(lower)alkoxyborane, di(lower)alkylborane,



and the other is halogen or $-\text{OSO}_2(\text{C}_q\text{F}_{2q+1})-$ wherein q is an integer of 0 to 4, or reacting a compound of the formula (II'):

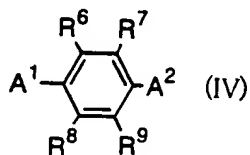


with a compound of the formula (III'):

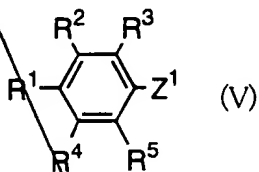


wherein, in the formulas (II') and (III'), $\text{R}^1\text{-R}^{13}$, X and Y are the same as defined [in claim 7] above and A and Z are the same as defined in the above formulas (II) and (III).

Claim 16 (Amended) The process for producing the compound of the formula (I''') according to claim 15, pharmaceutically acceptable salt or hydrate thereof [according to claim 15] comprising the reaction of a compound of the formula (IV):



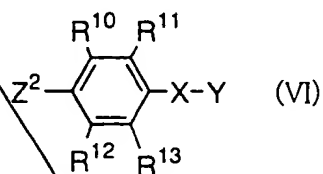
with a compound of the formula (V):



wherein, in the formulas (IV) and (V), R¹, R², R³, R⁴, R⁵, R⁶, R⁷, R⁸
and R⁹ are each independently hydrogen, hydroxy, halogen, carboxy,
optionally substituted lower alkyl, optionally substituted lower
alkoxy, optionally substituted lower alkenyl, optionally
substituted lower alkenyloxy, optionally substituted lower
alkylthio, optionally substituted lower alkoxy, optionally
substituted acyloxy, optionally substituted lower alkylsulfonyl,
optionally substituted lower alkylsulfonyloxy, optionally
substituted lower alkylsulfinyl, nitro, cyano, formyl, optionally
substituted amino, optionally substituted carbamoyl, optionally

substituted sulfamoyl or optionally substituted heterocyclyl, Z¹
is defined the same as for Z [defined] in the formula (II) [in
claim 15], A¹ and A² are each independently defined the same as for
A [defined] in the formula (III) [in claim 15], and the reactivity
of A¹ is higher than or equal to that of A²,

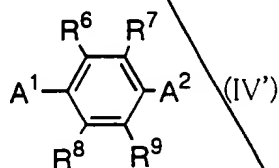
followed by the reaction with a compound of the formula (VI):



wherein R¹⁰-R¹³ are as defined for R¹-R⁹ above, X is -O-, -CH₂-,
NR¹⁴- wherein R¹⁴ is hydrogen, optionally substituted lower alkyl,
optionally substituted lower alkenyl or acetyl, or -S(O)_p- wherein
p is an integer of 0 to 2, Y is optionally substituted lower
alkyl, optionally substituted lower alkenyl, optionally
substituted lower alkynyl, optionally substituted acyl, optionally
substituted cycloalkyl, optionally substituted cycloalkenyl,
optionally substituted aryl or optionally substituted
heterocyclyl, and Y may optionally be substituted lower alkoxy
when X is -CH₂- and may optionally be substituted lower
alkoxycarbonyl, optionally substituted lower alkylsulfonyl or
optionally substituted arylsulfonyl when X is -O- or -NR¹⁴, R¹ and
R⁴, R¹ and R², R² and R³, R⁴ and R⁵, R⁶ and R⁷, R⁸ and R⁹, R¹⁰ and R¹¹,

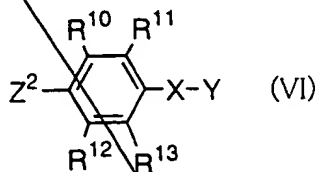
~~R^{12} and R^{13} , R^{11} and $-X-Y$, or R^{13} and $-X-Y$ taken together may form a 5- or 6-membered ring which may contain one or more of O, S or NR^{15} wherein R^{15} is hydrogen, optionally substituted lower alkyl, optionally substituted lower alkenyl, optionally substituted arylsulfonyl, and which may optionally be substituted, [X and Y are the same as defined in the formula (I) in claim 7] and Z^2 is the same as [Z defined in the above formula (II)] Z^1 above.~~

~~Claim 17 (Amended) The process for producing the compound of the formula (I''') according to claim 15, pharmaceutically acceptable salt or hydrate thereof [according to claim 15] comprising the reaction of a compound of the formula (IV'):~~



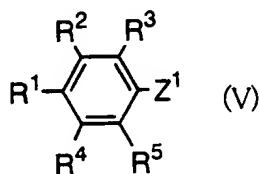
~~wherein, [R^6-R^9 is the same as defined in the formula (I) in claim 7,] R^6 , R^7 , R^8 and R^9 are each independently hydrogen, hydroxy, halogen, carboxy, optionally substituted lower alkyl, optionally substituted lower alkoxy, optionally substituted lower alkenyl, optionally substituted lower alkenyloxy, optionally substituted lower alkylthio, optionally substituted lower alkoxycarbonyl, optionally substituted acyloxy, optionally substituted lower~~

alkylsulfonyl, optionally substituted lower alkylsulfonyloxy,
optionally substituted lower alkylsulfinyl, nitro, cyano, formyl,
optionally substituted amino, optionally substituted carbamoyl,
optionally substituted sulfamoyl or optionally substituted
heterocyclyl. A¹ and A² are each independently defined the same as
 for A [defined] in the formula (III) [in claim 15], and the
 reactivity of A² is higher than or equal to that of A¹, with a
 compound of the formula (VI) [in claim 16]



wherein R¹⁰-R¹³ are as defined for R⁶-R⁹ above, X is -O-, -CH₂-,
 NR¹⁴- wherein R¹⁴ is hydrogen, optionally substituted lower alkyl,
 optionally substituted lower alkenyl or acetyl, or -S(O)_p- wherein
 p is an integer of 0 to 2, Y is optionally substituted lower
 alkyl, optionally substituted lower alkenyl, optionally
 substituted lower alkynyl, optionally substituted acyl, optionally

substituted cycloalkyl, optionally substituted cycloalkenyl,
 optionally substituted aryl or optionally substituted
 heterocyclyl, and Y may optionally be substituted lower alkoxy
 when X is $-\text{CH}_2-$ and may optionally be substituted lower
 alkoxy carbonyl, optionally substituted lower alkylsulfonyl or
 optionally substituted arylsulfonyl when X is $-\text{O}-$ or $-\text{NR}^{14}$, R^1 and
 R^4 , R^1 and R^2 , R^2 and R^3 , R^4 and R^5 , R^6 and R^7 , R^8 and R^9 , R^{10} and R^{11} ,
 R^{12} and R^{13} , R^{11} and $-\text{X}-\text{Y}$, or R^{13} and $-\text{X}-\text{Y}$ taken together may form a
 5- or 6-membered ring which may contain one or more of O, S or NR^{15}
 wherein R^{15} is hydrogen, optionally substituted lower alkyl,
 optionally substituted lower alkenyl, optionally substituted
 arylsulfonyl, and which may optionally be substituted, and Z^2 is
 defined the same as for Z in formula (II), followed by the
 reaction with a compound of the formula (V) [in claim 16.]



wherein R^1 - R^5 are as defined for R^6 - R^9 above, Z^1 is defined the same
 as for Z in the formula (II).

Please cancel claims 23-25 without prejudice or disclaimer to the subject matter contained therein.

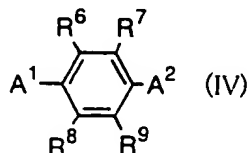
Please add the following new claims:

26. The selective suppressor of the IgE production claimed in claim 4 which suppresses infiltration of an inflammatory cell to tissue.

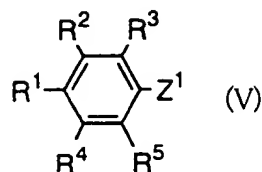
27. The selective suppressor of the IgE production claimed in claim 26 wherein the inflammatory cell is an eosinophil and/or a netrophile.

28. A pharmaceutical composition comprising the compound, pharmaceutically acceptable salt, hydrate or prodrug thereof claimed in claims 18, 19, 20, 21 or 22.

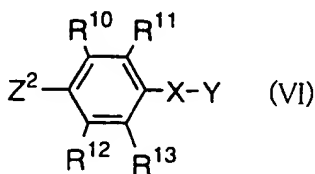
29. A process for producing a compound of the formula (I) according to claims 18, 19, 20, 21 or 22, pharmaceutically acceptable salt or hydrate thereof comprising reacting a compound of the formula (IV)



with a compound of the formula (V):



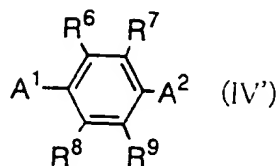
wherein, in the formulas (IV) and (V), R^1 , R^2 , R^3 , R^4 , R^5 , R^6 , R^7 , R^8 and R^9 are each independently hydrogen, hydroxy, halogen, carboxy, optionally substituted lower alkyl, optionally substituted lower alkoxy, optionally substituted lower alkenyl, optionally substituted lower alkenyloxy, optionally substituted lower alkylthio, optionally substituted lower alkoxycarbonyl, optionally substituted acyloxy, optionally substituted lower alkylsulfonyl, optionally substituted lower alkylsulfonyloxy, optionally substituted lower alkylsulfinyl, nitro, cyano, formyl, optionally substituted amino, optionally substituted carbamoyl, optionally substituted sulfamoyl or optionally substituted heterocyclyl, Z^1 is defined the same as for Z in the formula (II), A^1 and A^2 are each independently defined the same as for A in the formula (III), and the reactivity of A^1 is higher than or equal to that of A^2 , followed by the reaction with a compound of the formula (VI):



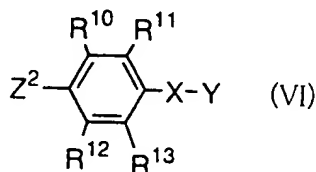
-NR¹⁴- wherein R¹⁴ is hydrogen, optionally substituted lower alkyl, optionally substituted lower alkenyl or acetyl, or -S(O)_p- wherein p is an integer of 0 to 2, Y is optionally substituted lower alkyl, optionally substituted lower alkenyl, optionally substituted lower alkynyl, optionally substituted acyl, optionally substituted cycloalkyl, optionally substituted cycloalkenyl, optionally substituted aryl or optionally substituted heterocyclyl, and Y may optionally be substituted lower alkoxy when X is -CH₂- and may optionally be substituted lower alkoxy carbonyl, optionally substituted lower alkylsulfonyl or optionally substituted arylsulfonyl when X is -O- or -NR¹⁴, R¹ and R⁴, R¹ and R², R² and R³, R⁴ and R⁵, R⁶ and R⁷, R⁸ and R⁹, R¹⁰ and R¹¹, R¹² and R¹³, R¹¹ and -X-Y, or R¹³ and -X-Y taken together may form a 5- or 6-membered ring which may contain one or more of O, S or NR¹⁵ wherein R¹⁵ is hydrogen, optionally substituted lower alkyl, optionally substituted lower alkenyl, optionally substituted arylsulfonyl, and which may optionally be substituted, and Z² is the same as Z¹ above.

30. A process for producing a compound of the formula (I), according to claims 18, 19, 20, 21 or 22 pharmaceutically acceptable salt or hydrate thereof comprising reacting a compound of the formula (IV')

acceptable salt or hydrate thereof comprising reacting a compound of the formula (IV')



wherein, R⁶, R⁷, R⁸ and R⁹ are each independently hydrogen, hydroxy, halogen, carboxy, optionally substituted lower alkyl, optionally substituted lower alkoxy, optionally substituted lower alkenyl, optionally substituted lower alkenyloxy, optionally substituted lower alkylthio, optionally substituted lower alkoxy carbonyl, optionally substituted acyloxy, optionally substituted lower alkylsulfonyl, optionally substituted lower alkylsulfonyloxy, optionally substituted lower alkylsulfinyl, nitro, cyano, formyl, optionally substituted amino, optionally substituted carbamoyl, optionally substituted sulfamoyl or optionally substituted heterocyclyl, A¹ and A² are each independently defined the same as for A in the formula (III), and the reactivity of A² is higher than or equal to that of A¹, with a compound of the formula (VI)



wherein R^{10} - R^{13} are as defined for R^6 - R^9 above, X is -O-, -CH₂-, -NR¹⁴- wherein R¹⁴ is hydrogen, optionally substituted lower alkyl, optionally substituted lower alkenyl or acetyl, or -S(O)_p- wherein p is an integer of 0 to 2, Y is optionally substituted lower alkyl, optionally substituted lower alkenyl, optionally substituted lower alkynyl, optionally substituted acyl, optionally substituted cycloalkyl, optionally substituted cycloalkenyl, optionally substituted aryl or optionally substituted heterocyclyl, and Y may optionally be substituted lower alkoxy when X is -CH₂- and may optionally be substituted lower alkoxy carbonyl, optionally substituted lower alkylsulfonyl or optionally substituted arylsulfonyl when X is -O- or -NR¹⁴, R¹ and R⁴, R¹ and R², R² and R³, R⁴ and R⁵, R⁶ and R⁷, R⁸ and R⁹, R¹⁰ and R¹¹, R¹² and R¹³, R¹¹ and -X-Y, or R¹³ and -X-Y taken together may form a 5- or 6-membered ring which may contain one or more of O, S or NR¹⁵ wherein R¹⁵ is hydrogen, optionally substituted lower alkyl, optionally substituted lower alkenyl, optionally substituted arylsulfonyl, and which may optionally be substituted, and Z² is defined the same as for Z in formula (II), followed by the reaction with a compound of the formula (V)

